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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
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EXAMINER

ANGELL, JON E

ART UNIT	PAPER NUMBER
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1635

DATE MAILED: 03/29/2002

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Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary

Application No.

09/763,682

Applicant(s)

BJERKVIG, ROLF

Examiner

J. Eric Angell

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-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133).
- Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☐ Responsive to communication(s) filed on ____.
- 2a) ☐ This action is **FINAL**. 2b) ☒ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 12-28 is/are pending in the application.
- 4a) Of the above claim(s) 23 is/are withdrawn from consideration.
- 5) ☐ Claim(s) ____ is/are allowed.
- 6) ☒ Claim(s) 12-22 and 24-28 is/are rejected.
- 7) ☐ Claim(s) ____ is/are objected to.
- 8) ☐ Claim(s) ____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on ____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
- Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
- 11) ☐ The proposed drawing correction filed on ____ is: a) ☐ approved b) ☐ disapproved by the Examiner.
- If approved, corrected drawings are required in reply to this Office action.
- 12) ☐ The oath or declaration is objected to by the Examiner.

Priority under 35 U.S.C. §§ 119 and 120

- 13) ☒ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☒ All b) ☐ Some * c) ☐ None of:
1. ☒ Certified copies of the priority documents have been received.
2. ☐ Certified copies of the priority documents have been received in Application No. ____.
3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).
- * See the attached detailed Office action for a list of the certified copies not received.
- 14) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. § 119(e) (to a provisional application).
- a) ☐ The translation of the foreign language provisional application has been received.
- 15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121.

Attachment(s)

- 1) ☒ Notice of References Cited (PTO-892)
- 2) ☐ Notice of Draftsperson's Patent Drawing Review (PTO-948)
- 3) ☐ Information Disclosure Statement(s) (PTO-1449) Paper No(s) ____.
- 4) ☐ Interview Summary (PTO-413) Paper No(s) ____.
- 5) ☐ Notice of Informal Patent Application (PTO-152)
- 6) ☐ Other:

DETAILED ACTION

Claims 12-28 are pending in the application.

Priority

1. Receipt is acknowledged of papers submitted under 35 U.S.C. 119(a)-(d), which papers have been placed of record in the file.

Election/Restrictions

2. Applicant's election without traverse of Group I, and the species of "a molecule that is capable of affecting tumor neovascularization" in Paper No. 8 is acknowledged. Claim 23 is drawn to a non-elected group and is withdrawn from consideration.

Claim Rejections - 35 USC § 112

3. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

4. Claims 12-22 and 24-28 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor(s), at the time the application was filed, had possession of the claimed invention.

The current claims are drawn to a composition comprising an encapsulated producer cell capable of producing a genus of any molecule that is an inhibitor of the growth of a CNS tumor

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that affects neovascularization. This large genus is represented in the specification by only a single species (endostatin). Thus, applicant has express possession of only 1 species in a genus which comprises a vast amount different possibilities (i.e. any molecule that inhibits CNS tumor growth and affects neovascularization). The written description guidelines note regarding such genus/species situations that "Satisfactory disclosure of a ``representative number" depends on whether one of skill in the art would recognize that the applicant was in possession of the necessary common attributes or features of the elements possessed by the members of the genus in view of the species disclosed." (See: Federal Register: December 21, 1999 (Volume 64, Number 244), revised guidelines for written description.) Here, there is only one element disclosed, therefore there are no predictable structures or attributes of the entire genus disclosed. There are no particular domains and no structural limitations or requirements which provide guidance on the identification of the CNS tumor growth inhibitors that affect neovascularization which meet these functional limitations provided.

It is noted that in Fiers v. Sugano (25 USPQ2d, 1601), the Fed. Cir. concluded that "...if inventor is unable to envision detailed chemical structure of DNA sequence coding for specific protein, as well as method of obtaining it, then conception is not achieved until reduction to practice has occurred, that is, until after gene has been isolated...conception of any chemical substance, requires definition of that substance other than by its functional utility."

In the instant application, only the CNS tumor growth inhibitor affecting neovascularization described is endostatin.

In Vas-Cath Inc. v. Mahurkar (19 USPQ2d 1111, CAFC 1991), it was concluded that:

"...applicant must also convey, with reasonable clarity to those skilled in art, that applicant, as of filing date sought, was in possession of invention, with invention being, for purposes of "written description" inquiry, whatever is presently claimed."

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In the application at the time of filing, there is no record or description which would demonstrate conception or written description of any other CNS tumor growth inhibitor that affects neovascularization.

5. Claims 12-22 and 24-28 rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for an encapsulated producer cell that expresses an inhibitor of the growth of a CNS tumor and affects neovascularization in rats, does not reasonably provide enablement for an encapsulated producer cell that expresses an inhibitor of the growth of a CNS tumor and affects neovascularization in any animal. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make/use the invention commensurate in scope with these claims.

Factors to be considered in determining whether a disclosure meets the enablement requirement of 35 USC 112, first paragraph, have been described by the court in *In re Wands*, 8 USPQ2d 1400 (CA FC 1988).

Wands states on page 1404,

“Factors to be considered in determining whether a disclosure would require undue experimentation have been summarized by the board in *Ex parte Forman*. They include (1) the quantity of experimentation necessary, (2) the amount of direction or guidance presented, (3) the presence or absence of working examples, (4) the nature of the invention, (5) the state of the prior art, (6) the relative skill of those in the art, (7) the predictability or unpredictability of the art, and (8) the breadth of the claims.”

The nature of the invention

The instant claims are drawn to a composition comprising a cell “capable” of expressing a molecule inhibiting growth of a CNS tumor, methods for making the composition and methods

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of using the composition. Therefore, the nature of the Invention encompasses gene therapy as well as cell encapsulation technology.

The breadth of the claims

The breadth of the claims is very broad. For instance the claims encompass a cell that is capable of expressing any molecule that inhibit the growth of a CNS tumor (including ones that affect neovascularization) including angiostatin, endostatin, any apoptotic inducing molecule and functional fragments thereof (just to name a few). There is no limitation in the claim of the type of molecule used to express the therapeutic agent; therefore the claims encompass the use of any vector including viral and non-viral vectors. Furthermore, the treatment claims encompass treating any CNS tumor in any species of animal, including humans.

The unpredictability of the art and the state of the prior art

The current relevant art also considers encapsulated cell technology for treatment of CNS disorders to be unpredictable. For instance, Visted et al. (Neuro-Oncology, July 2001, p. 201-210) recognizes many of the problems related to cell encapsulation technology. Visted et al. teaches that (1) “gene therapy using viral vectors has to date failed to reveal its definitive clinical usefulness” (p. 201, first paragraph); (2) the M component of the alginate used to encapsulate the cells “has immunogenic properties and it may evoke immune reactions when it is present at high concentrations (more than 85%) in the alginate” (see p. 202, under Alginate); (3) “despite promising results reported in several animal experiments, limited graft survival may occur. This is attributed to host immune reactions against the implant” (see p.204 under Biocompatibility); (4) “microcapsule graft failure is often associated with fibrotic outgrowth of the capsules” (see p. 205, first paragraph); (5) “several cases report that the host produces immunoglobulins against

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the encapsulated material as well as the secreted recombinant proteins” (see p. 205, under Reaction of the Recipient Against the Encapsulated Cells); (6) “The host’s tolerance to xenografts of encapsulated biomaterial also appears to vary between species. Therefore, if specific microcapsules are well tolerated in small animals, testing in large animals is a prerequisite before clinical application.” (see p. 205, under Reaction of the Recipient Against the Encapsulated Cells); (7) “only limited information is available on the parenchymal reaction to microcapsules. The alginates, with or without encapsulated producer cells, could theoretically elicit a brisk glial reaction that could abolish any therapeutic benefits” (see p. 205, under Reaction of the Recipient Against the Encapsulated Cells); (8) “There is also limited information available on alginate toxicity/reactivity in the brain in large animal models” (see p. 205, under Reaction of the Recipient Against the Encapsulated Cells); and (9) “Encapsulated cells, however, still have not been applied in any trial that included patients with malignant gliomas” (see p. 207, third paragraph).

Working Examples and Guidance in the Specification

The specification has only one working example of a treatment of a CNS tumor using an encapsulated cell expressing a molecule that affects neovascularization. The disclosed example is the treatment of a rat using an encapsulated cell expressing endostatin. The specification does not provide teachings sufficient to overcome doubts raised in the art with regards to methods of treatment of any animal other than a rat, or the use of any CNS tumor growth inhibitor other than endostatin. It would essentially be a trial and error process to make and use the diverse species of CNS growth inhibitors encompassed by the claims. It is further not predictable that the

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claimed treatment method would effectively achieve any therapeutic benefit in any animal other than a rat.

Quantity of Experimentation

The quantity of experimentation in this area is extremely large since determination of the efficacy of the encapsulated cell therapy would require, initially, large animal studies before the clinical trials (as mentioned by Visted et al.). After experimentation in the large animal model(s), the efficacy of the treatment would have to be tested in human subjects. This would require years of inventive effort, with each of the many intervening steps, upon effective reduction to practice, not providing any guarantee of success in the succeeding steps. Furthermore, all of the different possible growth inhibitors would have to be tested.

Level of the skill in the art

The level of the skill in the art is deemed to be high.

Conclusion

Considering the high degree of unpredictability of encapsulated cell technology recognized in the art, the breadth of the claims, the lack of working examples and guidance in the specification; and the high degree of skill required, it is concluded that the amount of experimentation required to perform the broadly claimed method is undue.

6. The following is a quotation of the second paragraph of 35 U.S.C. 112:

The specification shall conclude with one or more claims particularly pointing out and distinctly claiming the subject matter which the applicant regards as his invention.

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7. Claims 12-22 and 24-28 are rejected under 35 U.S.C. 112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which applicant regards as the invention.

Claim 12 recites the phrase, “ a producer cell **capable** of expressing a molecule... the cell being encapsulated in a matrix...” (emphasis added for clarity). The term “capable” renders the claim vague and indefinite because it means that the encapsulated cell may or may not express the molecule of interest, only that it is capable of expressing the molecule. For example, an encapsulated cell that does not express a particular molecule is still capable of expressing the molecule because it is possible to transfect/tranform the cell with an expression plasmid that expresses the molecule of interest. Therefore any encapsulated cell is capable of expressing a molecule of interest (such as a inhibitor of tumor growth).

Claims 12-22 and 24-28 depend on claim 12 and are therefore rejected for the same reason.

Claim Rejections - 35 USC § 102

8. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless –

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

9. When the claims are read as a cell that does not express a molecule that inhibits tumor growth, only one that is **capable** of expressing the molecule, claims 12-16, 18, 20 and 26 are

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rejected under 35 U.S.C. 102(b) as being anticipated by Skjak-Braek et al. (U.S. Patent 5,459,054).

Skjak-Braek et al. teaches teaches a composition comprising a producer cell which does not express a molecule that inhibits tumor growth, but is capable of expressing the molecule wherein the producer cell is encapsulated in a matrix that comprises an immunoisolating alginate having a G content of above 15%, above 50%, 60-80%, and 80-100%, wherein the producer cell is encapsulated in a bead, wherein the alginate is substantially pure of endotoxin, (see abstract; col. 4, lines 44-67; col. 7, lines 15-18; and Example 7).

Skjak-Braek et al. also teaches that the encapsulated cells are living cells (col. 4, lines 7-11) which are naturally occurring or genetically engineered prokaryotic or eukaryotic cells (see col. 4, lines 53-57).

Claim Rejections - 35 USC § 103

10. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

11. When the claims are read as a cell that does express a molecule that inhibits tumor growth, claims 12-16, 18, 20, 22 and 26 are rejected under 35 U.S.C. 103(a) as being unpatentable over Skjak-Braek et al. (U.S. Patent 5,459,054) in view of O'Reilly et al. (Cell, Vol. 88:277-285; Jan. 24, 1997).

Skjak-Braek et al. teaches a composition comprising a producer cell which does not express a molecule that inhibits tumor growth, but is capable of expressing the molecule wherein the producer cell is encapsulated in a matrix that comprises an immunoisolating alginate having a G content of above 15%, above 50%, 60-80%, and 80-100%, wherein the producer cell is encapsulated in a bead, wherein the alginate is substantially pure of endotoxin, (see abstract; col. 4, lines 44-67; col. 7, lines 15-18; and Example 7).

Skjak-Braek et al. also teaches that the encapsulated cells are living cells (col. 4, lines 7-11) which are naturally occurring or genetically engineered prokaryotic or eukaryotic cells (see col. 4, lines 53-57), and that the encapsulated cells “can be implanted or transplanted in vivo into mammals without inducing any substantial immunogenic reaction or fibroblast formation” and can be used “as a drug or biological material delivery system.” (See col. 4 lines 44-58).

Skjak-Braek et al. does not teach that the cell expresses a tumor growth suppressor molecule or that the molecule affects neovascularization (such as endostatin).

O'Reilly et al. teaches an inhibitor of angiogenesis and tumor growth known as endostatin (see abstract, fig. 4 and fig. 5). O'Reilly also teaches recombinant cells that express endostatin (see p. 284, left column).

It would have been prima facie obvious at the time of invention to combine the teachings of Skjak-Braek et al. and O'Reilly et al. and use the method cell encapsulation taught by Skjak-Braek et al. to encapsulate a cell which expresses endostatin with a reasonable expectation of success.

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The motivation to do so would have been to design a drug delivery system for delivering recombinant endostatin (expressed by an encapsulated cell that does not does not induce any substantial immunogenic reaction) to a subject in order to inhibit tumor growth.

Conclusion

No claim is allowed.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to J. Eric Angell whose telephone number is (703) 605-1165. The examiner can normally be reached on M-F (8:00-4:30).

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, John L. LeGuyader can be reached on (703) 308-0447. The fax phone numbers for the organization where this application or proceeding is assigned are (703) 308-4242 for regular communications and (703) 308-4242 for After Final communications.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist whose telephone number is (703) 308-0196.

J. Eric Angell, Ph.D.
March 25, 2002



**JEFFREY FREDMAN
PRIMARY EXAMINER**